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Amendments to the Claims:

This listing of claims replaces all prior versions and listings of claims in the application:

<u>Listing of Claims</u>:

- 1. (Original) An antibody recognizing both an enzyme and a substrate thereof, wherein said antibody is a bispecific antibody which functionally substitutes for a cofactor that enhances the enzymatic reaction.
- 2. (Original) The antibody according to claim 1, wherein said enzyme is a proteolytic enzyme.
- 3. (Original) The antibody according to claim 2, wherein said proteolytic enzyme, substrate, and cofactor are blood coagulation/fibrinolysis-associated factors.
- 4. (Original) The antibody according to claim 3, wherein the enzyme of a blood coagulation/fibrinolysis- associated factor is blood coagulation factor IX and/or activated blood coagulation factor IX; the substrate is blood coagulation factor X; and the cofactor is blood coagulation factor VIII and/or activated blood coagulation factor VIII.
- 5. (Currently Amended) The antibody according to claim 1 any one of claims 1 to 4, wherein said antibody comprises a complementarity determining region comprising the amino acid sequence of anti-blood coagulation factor IX/IXa antibody CDR3 of the following (a1) or (a2) or a complementarity determining region functionally equivalent thereto, and a complementarity determining region comprising the amino acid sequence of anti-blood coagulation factor X antibody CDR3 described in any one of the following (b1) to (b9) or a complementarity determining region functionally equivalent thereto:
 - (a1) H chain CDR 3 amino acid sequence described in SEQ ID NO: 16;

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(a2) H chain CDR 3 amino acid sequence described in SEQ ID NO: 20;

(b1) H chain CDR 3 amino acid sequence described in SEQ ID NO: 24;

(b2) H chain CDR 3 amino acid sequence described in SEQ ID NO: 28;

(b3) H chain CDR 3 amino acid sequence described in SEQ ID NO: 32;

(b4) H chain CDR 3 amino acid sequence described in SEQ ID NO: 36;

(b5) H chain CDR 3 amino acid sequence described in SEQ ID NO: 40;

(b6) H chain CDR 3 amino acid sequence described in SEQ ID NO: 44;

(b7) H chain CDR 3 amino acid sequence described in SEQ ID NO: 48;

(b8) H chain CDR 3 amino acid sequence described in SEQ ID NO: 52;

(b9) H chain CDR 3 amino acid sequence described in SEQ ID NO: 56.

- 6. (Currently Amended) The antibody according to <u>claim 1</u> any one of claims 1 to 4, wherein said antibody comprises a complementarity determining region comprising the amino acid sequence of anti-blood coagulation factor IX/IXa antibody CDR of the following (a1) or (a2) or a complementarity determining region functionally equivalent thereto, and a complementarity determining region comprising the amino acid sequence of anti-blood coagulation factor X antibody CDR described in any one of the following (b1) to (b9) or a complementarity determining region functionally equivalent thereto:
- (a1) H chain CDR 1, 2, and 3 amino acid sequences described in SEQ ID NOs: 14, 15, and 16, respectively;
- (a2) H chain CDR 1, 2, and 3 amino acid sequences described in SEQ ID NOs: 18, 19, and 20, respectively;
- (b1) H chain CDR 1, 2, and 3 amino acid sequences described in SEQ ID NOs: 22, 23, and 24, respectively;
- (b2) H chain CDR 1, 2, and 3 amino acid sequences described in SEQ ID NOs: 26, 27, and 28, respectively;
- (b3) H chain CDR 1, 2, and 3 amino acid sequences described in SEQ ID NOs: 30, 31, and 32, respectively;

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(b4) H chain CDR 1, 2, and 3 amino acid sequences described in SEQ ID NOs: 34, 35, and 36, respectively;

- (b5) H chain CDR 1, 2, and 3 amino acid sequences described in SEQ ID NOs: 38, 39, and 40, respectively;
- (b6) H chain CDR 1, 2, and 3 amino acid sequences described in SEQ ID NOs: 42, 43, and 44, respectively;
- (b7) H chain CDR 1, 2, and 3 amino acid sequences described in SEQ ID NOs: 46, 47, and 48, respectively;
- (b8) H chain CDR 1, 2, and 3 amino acid sequences described in SEQ ID NOs: 50, 51, and 52, respectively;
- (b9) H chain CDR 1, 2, and 3 amino acid sequences described in SEQ ID NOs: 54, 55, and 56, respectively.
- 7. (Currently Amended) A composition comprising the antibody according to <u>claim 1</u> any one of claims 1 to 6 and a pharmaceutically acceptable carrier.
- 8. (Original) The composition according to claim 7, wherein said composition is a pharmaceutical composition used for preventing and/or treating bleeding, disorder accompanied by bleeding, or disorder caused by bleeding.
- 9. (Original) The composition according to claim 8, wherein the bleeding, disorder accompanied by bleeding, or disorder caused by bleeding is a disorder that arises and/or progresses as a result of an activity decrease or deficiency of blood coagulation factor VIII and/or activated blood coagulation factor VIII.
- 10. (Original) The composition according to claim 9, wherein the disorder that arises and/or progresses as a result of an activity decrease or deficiency of blood coagulation factor VIII and/or activated blood coagulation factor VIII is hemophilia A.

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11. (Original) The composition according to claim 9, wherein the disorder that arises and/or progresses as a result of an activity decrease or deficiency of blood coagulation factor VIII and/or activated blood coagulation factor VIII is a disorder in which an inhibitor against blood coagulation factor VIII and/or activated blood coagulation factor VIII is generated.

- 12. (Original) The composition according to claim 9, wherein the disorder that arises and/or progresses as a result of an activity decrease or deficiency of blood coagulation factor VIII and/or activated blood coagulation factor VIII is acquired hemophilia.
- 13. (Original) The composition according to claim 9, wherein the disorder that arises and/or progresses as a result of an activity decrease of blood coagulation factor VIII and/or activated blood coagulation factor VIII is von Willebrand's disease.
- 14. (Currently Amended) A method for preventing and/or treating bleeding, disorder accompanied by bleeding, or disorder caused by bleeding, wherein said method comprises the step of administering the antibody according to <u>claim 1</u> any one of claims 1 to 6, or the <u>composition according to any one of claims 7 to 13</u>.
- 15. (Currently Amended) Use of the antibody according to <u>claim 1</u> any one of claims 1 to 6 for the preparation of a pharmaceutical composition preparing the composition according to any one of claims 7 to 13.
- 16. (Currently Amended) A kit <u>used in a method of preventing and/or treating bleeding</u>, <u>disorder accompanied by bleeding</u>, <u>or disorder caused by bleeding used in the method of preventing and/or treating disorders according to claim 14</u>, wherein said kit comprises at least the antibody according to <u>claim 1</u> any one of claims 1 to 6 or the composition according to claim 7.

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17. (Currently Amended) A method of preventing and/or treating bleeding, disorder accompanied by bleeding, or disorder caused by bleeding, wherein said method comprises the step of administering the antibody according to claim 4 any one of claims 4 to 6 or the composition according to any one of claims 7 to 13 in combination with blood coagulation factor VIII.

- 18. (Currently Amended) A kit used in [[the]] a method of preventing and/or treating bleeding, disorder accompanied by bleeding, or disorder caused by bleeding according to claim 17, wherein said kit comprises at least the antibody according to claim 4 any one of claims 4 to 6, or the composition according to claim 7, and blood coagulation factor VIII.
- 19. (New) A method for preventing and/or treating bleeding, disorder accompanied by bleeding, or disorder caused by bleeding, wherein said method comprises the step of administering the composition according to claim 7.